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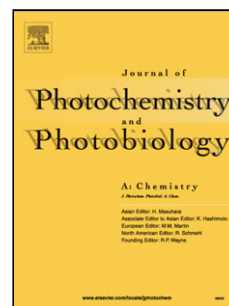
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## Improvement of sol-gel prepared tungsten trioxide photoanodes upon doping with ytterbium

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### **HIGHLIGHTS**

- A sol-gel method for the preparation of Yb-doped  $\text{WO}_3$  photoanodes is presented.
- Ytterbium modification induces an increase in the photoresponse of  $\text{WO}_3$  electrodes.
- Yb changes  $\text{WO}_3$  morphology and surface charge without increasing carrier density.

### **Abstract**

The development of efficient and stable photoelectrodes keeps on being critical in the development of practical water splitting devices. Sol-gel synthesized tungsten trioxide thin film electrodes modified with ytterbium have been prepared on conducting glass substrates. All the obtained electrodes were found to be composed of monoclinic  $\text{WO}_3$ ,

which is the photoactive crystal phase of  $\text{WO}_3$  for water oxidation. The photoelectrochemical behavior of both  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  nanostructured electrodes in 0.1 M  $\text{HClO}_4$  is apparently characterized by a low degree of recombination. Ytterbium addition to tungsten trioxide improves its photoelectrochemical response as larger photocurrents are obtained without a significant change of the onset potential. This behavior is attributed primarily to a surface effect rather than on bulk doping. The addition of  $\text{H}_2\text{O}_2$  to the aqueous electrolyte gives rise to both an increment in photocurrent and a shift of the photocurrent onset potential toward less positive values, which can be linked to the fact that hydrogen peroxide acts as an efficient hole scavenger. The implications of these findings in the mechanism of water photooxidation on  $\text{WO}_3$  are discussed.

**Keywords:** photoelectrochemical, tungsten trioxide, water splitting, ytterbium doping.

## 1. Introduction

Nowadays, the depletion of fossil energy resources is a serious problem which will be affecting modern life as we conceive it now. Finding carbon free energy sources is one of the most pressing challenges of humanity [1,2]. Furthermore, since Fujishima and Honda published the first photoelectrochemical (PEC) water splitting study in 1972 [3], hydrogen has been widely studied as a promising energy vector that can be produced in a renewable way. Taking into account that solar energy is an inexhaustible natural resource and that hydrogen is a potential candidate to be used as a clean fuel, solar water splitting seems to be one of the most convenient ways to harvest solar light and to convert it into chemical energy, hydrogen for instance [4-6].

Most of the PEC devices work in combination with photovoltaic elements (e.g., p-n junctions) and contain one photoelectrode (normally a photoanode) and one dark electrode [6]. Nevertheless, a more straightforward and potentially more efficient path to light-driven chemical transformations is to directly interface semiconductors and an appropriate electrolyte in a PEC tandem cell, where there is one photoanode and one photocathode [7-9]. If we focus on oxide electrodes, Sivula et al. [9] have shown that a bismuth vanadate-cuprous oxide tandem cell for overall solar water splitting can work without applying an external potential, although the stability of the oxide electrodes is limited.

Photoanodes used in PEC water splitting are n-type semiconductors that under illumination can photogenerate holes that oxidize water into  $O_2$ . They mainly include various binary and ternary oxides [5], for instance,  $\alpha\text{-Fe}_2\text{O}_3$  [10,11],  $\text{WO}_3$  [12-15],  $\text{ZnO}$  [16,17],  $\text{In}_2\text{O}_3$  [18],  $\text{SnO}_2$  [19],  $\text{TiO}_2$  [20-21],  $\text{BiVO}_4$  [22-24],  $\text{CuWO}_4$  [24,25],  $\text{Bi}_2\text{WO}_6$  [25], and bismuth molybdates ( $\text{Bi}_2\text{O}_3 \cdot n\text{MoO}_3$ ) [26]. Among them, tungsten oxide ( $\text{WO}_3$ ) has been extensively studied as a photoanode for use in solar water splitting since its valence band maximum position can provide sufficient overpotential for

photogenerated holes to oxidize water [27]. Furthermore,  $\text{WO}_3$  is not only inexpensive and nontoxic, but also it is one of the few oxides that are stable in acidic aqueous media, which opens up the possibility of preparing tandem cells based on acidic electrolytes [5,13,28].

The photooxidation of small organic molecules in competition with that of water has been studied as a way to understand the photoelectrochemical response of  $\text{WO}_3$  photoanodes and to assess their potential use in decontaminative photoelectrocatalysis [29,30]. It is remarkable that in all cases a very high conversion efficiency value was found. The oxidation of formic acid not only on electrodeposited, but also on doctor-blade deposited  $\text{WO}_3$  nanostructured thin film electrodes has been studied [13], being reported in both cases photocurrent doubling and a shift of the photocurrent onset potential toward less positive values. Remarkably, adding small organic molecules improves photoelectrochemical behavior as these species tend to be more easily oxidized than water molecules [13,31].

Other convenient ways for enhancing the photoelectrochemical performance of  $\text{WO}_3$  are modifying both morphology and/or composition (doping) [4,27]. Kalanur et al. [32], demonstrated that a significant increase in photocurrent is linked to the existence of direct pathways for electron transport such as those provided by vertically aligned nanorod arrays of  $\text{WO}_3$ . Regarding doping,  $\text{WO}_3$  has been doped with elements such as nitrogen [33,34], carbon [35], tellurium [36], cesium [37], and ytterbium [38]. Liew et al. [38] prepared Yb-doped  $\text{WO}_3$  by co-sputtering and found an improvement in photocurrent density, which was attributed to conductive carrier path and increased oxygen vacancies due to the substitution of  $\text{W}^{6+}$  by  $\text{Yb}^{3+}$  and electron enrichment by the  $4f^{13}$  orbital configuration of  $\text{Yb}^{3+}$ . Furthermore, they showed that Yb incorporation reduced charge transfer resistances and increased donor densities of  $\text{WO}_3$  photocatalysts. It is also worth noting that recent work in our laboratory has shown that Yb doping of hematite electrodes is effective for promoting photoactivity [39].

In this work, we report on the behavior of  $\text{WO}_3$  thin film electrodes prepared by a novel sol-gel procedure with a focus on their structural, morphological and photoelectrochemical properties. In addition,  $\text{WO}_3$  thin film electrodes were modified with ytterbium for the first time following a sol-gel procedure. It should be highlighted that our work is focused in ultrathin films since this avoids difficulties arising from composition variations throughout the thickness of the electrode as well as complexities derived from inhomogeneous illumination. This facilitates gaining physical insight on the dopant effect. In order to find the optimum amount of dopant for an enhanced photoelectrochemical behavior,  $\text{WO}_3$  electrodes were doped with five different atomic percentages of ytterbium. The photoelectrochemical response of, not only  $\text{WO}_3$  electrodes, but also  $\text{WO}_3\text{:Yb}$  in contact with a more effective hole scavenger than water has also been addressed. Important features of those electrodes such as photocurrent onset potential, photoactivity improvement and chemical stability are highlighted.

## 2. Experimental

$\text{WO}_3$  photoanodes were prepared on commercial fluorine-doped tin oxide (FTO) glass substrates by spin-coating followed by a two-step thermal treatment.  $\text{WCl}_6$  (150 mg, purity 99.99%+, Aldrich Chem. Co.) and polyethylene glycol (25 mg, purity 99%, Aldrich Chem. Co.) were dissolved in 1.25 mL ethanol and stirred for 4 hours [40]. Then, this precursor was spin-coated onto FTO at 1500 rpm for 15 s. The samples were dried, maintained at 40 °C in air for 30 minutes and then, annealed at 500 °C for 30 minutes at a ramp rate of 5 °C/min. On the other hand, to obtain  $\text{WO}_3$  doped with Yb (0.1, 0.2, 0.3, 0.4 y 0.5 %at Yb),  $\text{YbCl}_3 \cdot 6\text{H}_2\text{O}$  (purity 99.99%+, Aldrich Chem. Co.) was added to the precursor solution at the beginning of the synthetic procedure.

A Bruker D8-Advance X-ray diffractometer operating with Cu-K radiation at 40 kV and 40 mA was used to obtain the X-ray diffraction patterns. The composition of the electrodes was analyzed by X-ray photoelectron spectroscopy (XPS) using a Ka ThermoScientific spectrometer. The electrode morphology was studied with a Zeiss model MERLIN compact VP field emission scanning electron microscope (FE-SEM). The optical characterization of the electrodes was performed by solid-state UV-vis spectroscopy, using a Shimadzu UV-2401 PC spectrophotometer equipped with an integrating sphere. The UV-vis spectral data for the annealed films coated on transparent FTO conducting substrates were recorded in the absorbance mode.

Photoelectrochemical measurements were performed at room temperature using a three-electrode cell provided with a fused silica window and a computer-controlled Autolab PGSTAT30 potentiostat. In all cases, an Ag/AgCl/KCl(sat) electrode and a Pt wire were employed as a reference and counter electrode, respectively. All the potentials are referred to the Ag/AgCl/KCl(sat) electrode. Solutions of 0.1 M HClO<sub>4</sub> were used as working electrolytes. They were prepared with ultrapure water and N<sub>2</sub>-purged before the experiments. The illumination was carried from the electrolyte side (EE illumination) by employing a 300 W Xe-arc lamp (Thermo Oriel) equipped with a water filter. The light intensity was 0.120 mW·cm<sup>-2</sup> and it was measured by coupling an optical power meter (Thorlabs model PM100D) with a thermopile.

### **3. Results and discussions**

#### *3.1. Structural and physical characterization.*

To verify the chemical structure of the as-prepared thin films, they were analyzed by XRD. For WO<sub>3</sub> and WO<sub>3</sub>:Yb 0.2 at% samples, their XRD patterns shown in Fig. 1

indicate that the  $\text{WO}_3$  monoclinic phase (JCPDS 01-072-0677) is the only crystalline phase detected for both samples, which is the polymorph of  $\text{WO}_3$  more photoactive for water oxidation [32]. Therefore the introduction of small quantities of the dopant does not trigger any structural transformation.

Fig. 2 illustrates the morphology of the  $\text{WO}_3$  thin films with and without Yb. In both cases, nanocrystals are homogeneously distributed on the surface, being connected to one another. Grains have an average diameter close to 60 nm not only for both samples although it appears to be slightly smaller in the case of the  $\text{WO}_3\text{:Yb}$  sample (see histograms in the insets). Particle arrangement seems to be more uniform for  $\text{WO}_3\text{:Yb}$ , which can be attributed to a change of the sol-gel process included by the presence of Yb. In addition, the size distribution is more symmetrical and slightly narrower for Yb-doped samples.

In order to confirm that Yb is effectively introduced in the  $\text{WO}_3$  matrix, XPS analysis was performed, being the high resolution spectra for the W 4f, O 1s and Yb 4d regions displayed in Figure 3. The results point to the presence of Yb in the  $\text{WO}_3\text{:Yb}$  sample, which means that the incorporation of Yb is successfully performed. As expected for the small loading of Yb in the sample doped and intrinsic  $\text{WO}_3$  films show a similar W 4f spectra, which consists of two peaks centered at 35.46 eV and 37.58 eV associated with the spin-orbit doublet of W 4f<sub>7/2</sub> and W 4f<sub>5/2</sub> respectively. The O 1s spectra are also similar for both samples showing a peak at 530.28 eV typical of O-W bonding and a shoulder at 531.87 eV typical of O-C bonding coming from spurious impurities [38]. For the Yb 4d spectrum, a broad peak of low intensity at 185.1 eV can be assigned to Yb 4d<sub>5/2</sub> in good agreement with reported value of 185.0 eV for  $\text{Yb}_2\text{O}_3$  by Uwamino et al. [41]. This means that the environment for Yb in the  $\text{WO}_3$  lattice is similar to that found in  $\text{Yb}_2\text{O}_3$  (+3 oxidation state and coordination to oxygen). It is worth noting that even when 0.2 at% doped samples were prepared, a composition of 0.24 at% was reported



by XPS analysis, which suggests that there is a slight enrichment of Yb on the thin film surface.

The optical bandgap was obtained from optical absorption experiments. Fig. 4a shows the optical absorption spectra of  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  0.2 at% samples, which are very similar. However,  $\text{WO}_3\text{:Yb}$  samples have a slightly higher absorption background and there is a minor shift in the absorption edge. Figure 4b shows the Tauc plot of  $(Ah\nu)^{1/2}$  versus  $h\nu$  ( $A$  is the absorbance and  $h\nu$  the incident photon energy). Extrapolating the linear region, the value of  $E_g$  is estimated to be 2.65 eV for both pristine and doped  $\text{WO}_3$  samples, in agreement with the values between 2.6 and 2.8 eV reported in the literature [2,5].

### 3.2. Electrochemical characterization.

Cyclic voltammograms (CV) for  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  0.2 at% electrodes in the dark are shown in Fig. 5 in 0.1 M both in the dark (a) and under illumination (b). In the dark, the electrochemical response exhibits the existence of small capacitive currents in the low potential region (below 0.2 V), which tend to be higher when ytterbium is present. As these currents have demonstrated to be proportional to the interfacial area [21], it seems that the Yb-induced change could be mainly due to small changes in particle size and distribution as observed in Fig. 2. Importantly, the addition of Yb to the  $\text{WO}_3$  structure does not alter the shape of the voltammograms in the accumulation region, which point to the fact that doping does not alter in a significant way the conduction band/band gap states. In addition, this quasi-reversible process appears accompanied by blue coloration of the film at negative potentials. This process can be explained on the basis of proton intercalation leading to a colored tungsten bronze [Eq. (1)]:



Linear scan voltammograms performed under chopped illumination for  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  at 0.2 at% electrodes are shown in Fig. 5 (b). Upon illumination, anodic photocurrents appear with an onset potential at  $0.30 \text{ V}_{\text{Ag/AgCl}}$ , that is, at  $0.55 \text{ V}$  versus the reversible hydrogen electrode (RHE). In agreement with previous reports [13,42], the material behaves as an n-type semiconductor with a valence band located at around  $3.2 \text{ V}_{\text{RHE}}$  (by considering a bandgap value of  $2.65 \text{ eV}$  as determined above). Furthermore, there is an increase in the photocurrent density for  $\text{WO}_3\text{:Yb}$  0.2 at% samples.

The n-type behavior of the as-prepared electrodes was confirmed by constructing Mott-Schottky plots (Fig. 6, Fig. S1) at two different frequencies in the dark and under illumination in  $0.1 \text{ M HClO}_4$ . The slopes of the linear regions increase with frequency and for each frequency, the slope decreases in the order  $\text{WO}_3 < \text{WO}_3\text{:Yb}$ . This behavior in the slope might be directly related with a change not only in donor density but also in roughness of the electrode material, as it has been proposed by Gómez and coworkers [11]. In other words, since the real interfacial area for  $\text{WO}_3$  electrodes is much larger than the geometric one and, as long as characteristic dimension of the nano-objects is larger than that of the space charge region, the Mott-Schottky equation should be written as [Eq. (2)]:

$$\frac{A_{\text{real}}^2}{C_{\text{sc}}^2} = \frac{2}{e\epsilon_{\text{sc}}\epsilon_0 N_d} \left( E - E_{fb} - \frac{kT}{e} \right) \quad (2)$$

where  $C_{\text{sc}}$  is the capacitance of the space charge region,  $A_{\text{real}}$  is the real surface area, which takes into account the electrode roughness,  $\epsilon_{\text{sc}}$  is the semiconductor dielectric constant,  $\epsilon_0$  is the vacuum permittivity,  $N_d$  is the donor density,  $E$  is the applied potential, and  $E_{fb}$  is the flat band potential. Taking into account that the roughness factor ( $r$ ) can be expressed as [Eq. (3)]:

$$r = \frac{A_{\text{real}}}{A_{\text{geom}}} \quad (3)$$

where  $A_{geom}$  is the geometric (projected) area of the electrode, Equation 2 can be rewritten as [Eq. (4)]:

$$\frac{A_{geom}^2}{C_{SC}^2} = \frac{2}{e\epsilon_{sc}\epsilon_0 N_d r^2} \left( E - E_{fb} - \frac{kT}{e} \right) \quad (4)$$

To analyze the difference in the slopes, we must take into account that:

- The SEM analysis suggests that roughness factor is higher for WO<sub>3</sub>:Yb than for WO<sub>3</sub>, as the average particle size is slightly smaller for the Yb-containing sample.
- The capacity of the accumulation region for WO<sub>3</sub>:Yb is higher than for WO<sub>3</sub> (Fig. 5a), which is related with the real area of the electrodes, which means that roughness is higher for WO<sub>3</sub>:Yb than for WO<sub>3</sub>.

Therefore, if  $r_{WO_3:Yb} > r_{WO_3}$ , it must occur that  $N_{d,WO_3:Yb} < N_{d,WO_3}$ . This behavior might be related to a diminution of the density of oxygen vacancies upon addition of Yb [43]. The diminution in donor density is concomitant with an enhancement of the n-type behavior of the electrode. Such an apparent contradiction can be understood on the bases of an increase in electron mobility upon Yb doping even when their concentration decreases.

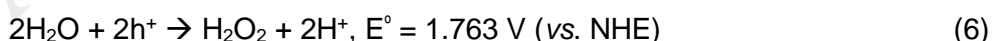
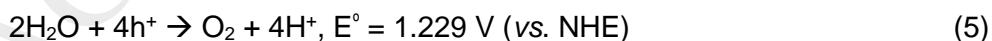
The flat band potential estimated from extrapolation of the intercept of the Mott-Schottky plot linear region with the x-axis, has approximately the same value (-0.05 V<sub>Ag/AgCl</sub>) for pure WO<sub>3</sub> and WO<sub>3</sub>:Yb electrodes. It is worth noting that this value is close to that of the photocurrent onset, indicating that surface recombination is limited in both cases. In agreement, the flat band potential does not significantly shift upon illumination, which means that the density of trapped holes under photostationary conditions when the illumination is intense does not induce an upward shift of the flat band potential.

The dependence of the photocurrent density on the atomic percent of Yb in the WO<sub>3</sub> electrodes is shown in Fig. 7 for two different values of the applied potential. Five

different amounts of atomic percentages (0.1, 0.2, 0.3, 0.4 and 0.5 at. %) were studied as to optimize the doping condition. As observed, the photocurrent is higher for the 0.2 at% sample not only at 0.6 V but also at 1.2 V. Thus, this is the adequate level of doping for a higher photoresponse. It should be emphasized that the observed non-linear dependence of the photoactivity on dopant concentration probably reflects the concurrence of several factors in determining the effect of Yb doping.

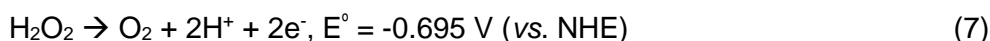
Fig. 8 shows the voltammetric behavior both in the dark and under illumination for pristine and Yb-modified WO<sub>3</sub> in the presence 0.1 M H<sub>2</sub>O<sub>2</sub>. The dark response of the WO<sub>3</sub> electrodes is characterized by the development of large cathodic currents associated to the reduction of H<sub>2</sub>O<sub>2</sub> whose onset coincides with that of the accumulation region. Upon illumination and irrespective of the presence of Yb, there is a shift of the onset potential toward less positive values and an increase in the photocurrent as observed by Gómez and coworkers [13] in the case of formic acid photooxidation. This is a consequence of adding H<sub>2</sub>O<sub>2</sub>, which can be linked to the fact that the hydrogen peroxide acts as an efficient hole scavenger. However, the increment in the photocurrent density of WO<sub>3</sub>:Yb at 0.2% atomic samples keeps on occurring at a level similar to that in the absence of H<sub>2</sub>O<sub>2</sub>. One should conclude that the improvement caused by doping with Yb is not directly linked to the charge transfer process, indirectly supporting the increase in electron mobility invoked above.

The efficiency of H<sub>2</sub>O<sub>2</sub> as a hole scavenger is quite evident. Water photooxidation could follow two possible pathways depicted in [Eq.(5)] and [Eq.(6)]:



In this context, [Eq.(5)], which does not consider direct formation of H<sub>2</sub>O<sub>2</sub>, is thermodynamically favorable, since the standard potential is significantly smaller than when its formation is required [Eq.(6)]. However, in a case when H<sub>2</sub>O<sub>2</sub> is already

present in the solution, its conversion to oxygen [Eq.(7)] becomes the primary reaction as the standard potential for this process is much lower than for the competitive process of water photooxidation according to the [Eq.(5)]. The latter illustrates the fact that  $\text{H}_2\text{O}_2$  can be readily oxidized even in the absence of a strong oxidizing agents (e.g. holes), and thus serve as a very efficient hole scavenger.



Based on these observations, it is plausible to assume that the formation of peroxo species is limiting the rate of water oxidation on  $\text{WO}_3$  photoanodes.

Finally, the pH effect is summarized in Fig. 9, which shows that the higher the pH value, the worst the photoresponse. It is worth noting that in the case of water photooxidation the  $\text{WO}_3\text{:Yb}$  electrode is much more sensitive to pH than the pristine  $\text{WO}_3$  electrode. This could be linked with the fact that there is Yb located at the surface, as demonstrated by XPS analysis, and the acid-base properties of the W-O and Yb-O species might be different, as well as, their isoelectric points (IEP), since IEP of  $\text{Yb}_2\text{O}_3$  is located at pH values around 7.5 – 8.5 while that of  $\text{WO}_3$  is located at pH values at around 1.5 – 2 [44]. Therefore, the change in the charge state of the surface (resulting from acid/base equilibria) would be the main factor responsible for the effects observed as a function of pH without the need of invoking Yb surface enrichment.

#### 4. Conclusions

In summary, we have shown that Yb modification of sol-gel synthesized tungsten trioxide improves its photoelectrochemical response as photocurrents increase while the onset potential remains virtually unaltered. Taking into account Mott-Schottky plots, this behavior is attributed to a surface effect and/or to an increase in electron mobility together with a slight diminution in particle size upon Yb modification. In addition, since

doping does not cause a significant alteration of the shape of the dark CVs in the accumulation region, no significant changes in the conduction band/band gap states seem to occur upon doping. Different doping levels were studied and the highest photocurrent was obtained for 0.2 at% Yb. The photoelectrochemical behavior of both  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  nanostructured electrodes in aqueous solutions of 0.1 M  $\text{HClO}_4$  is apparently characterized by a low degree of recombination. As a result, the addition of  $\text{H}_2\text{O}_2$  to the aqueous electrolyte only gives rise to a moderate increment in photocurrent and a slight shift of the photocurrent onset potential towards less positive values, despite  $\text{H}_2\text{O}_2$  being a very effective hole scavenger. Importantly, the addition of  $\text{H}_2\text{O}_2$  causes similar effects regardless the presence of Yb, meaning that factors different from the hole transfer process, such as charge carrier mobility play a crucial role in the Yb-induced enhancement.

## 5. Acknowledgements

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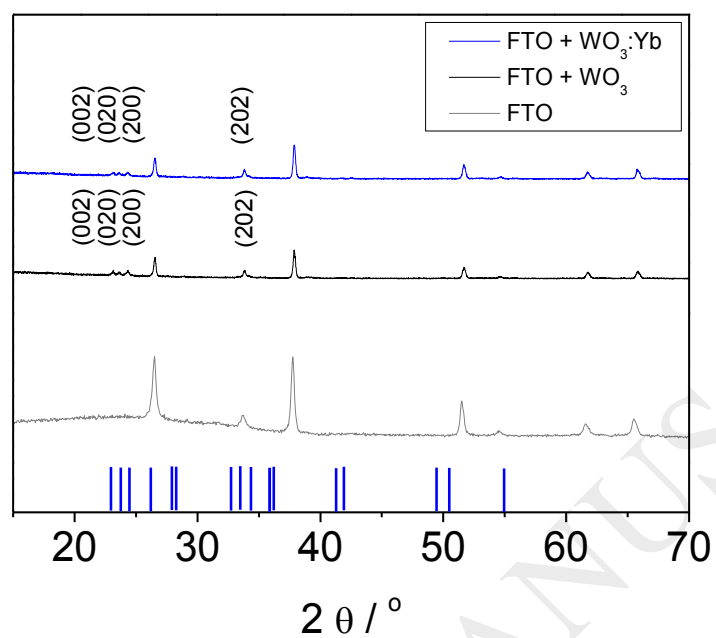


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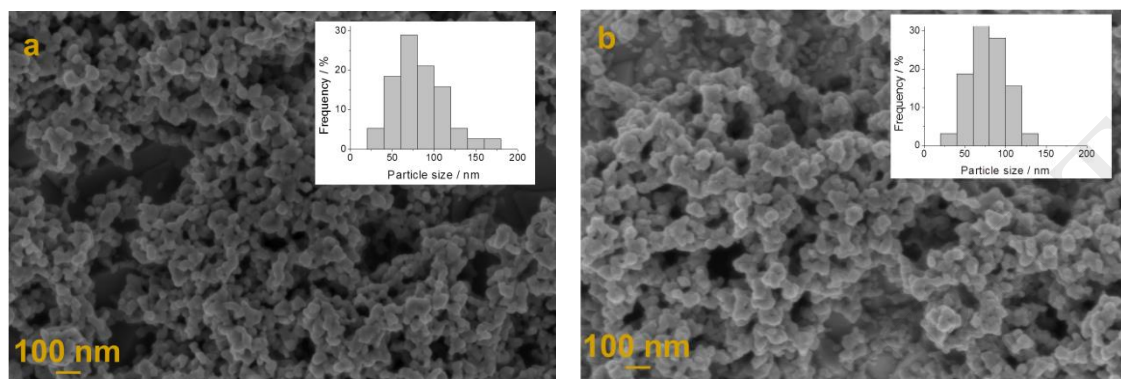
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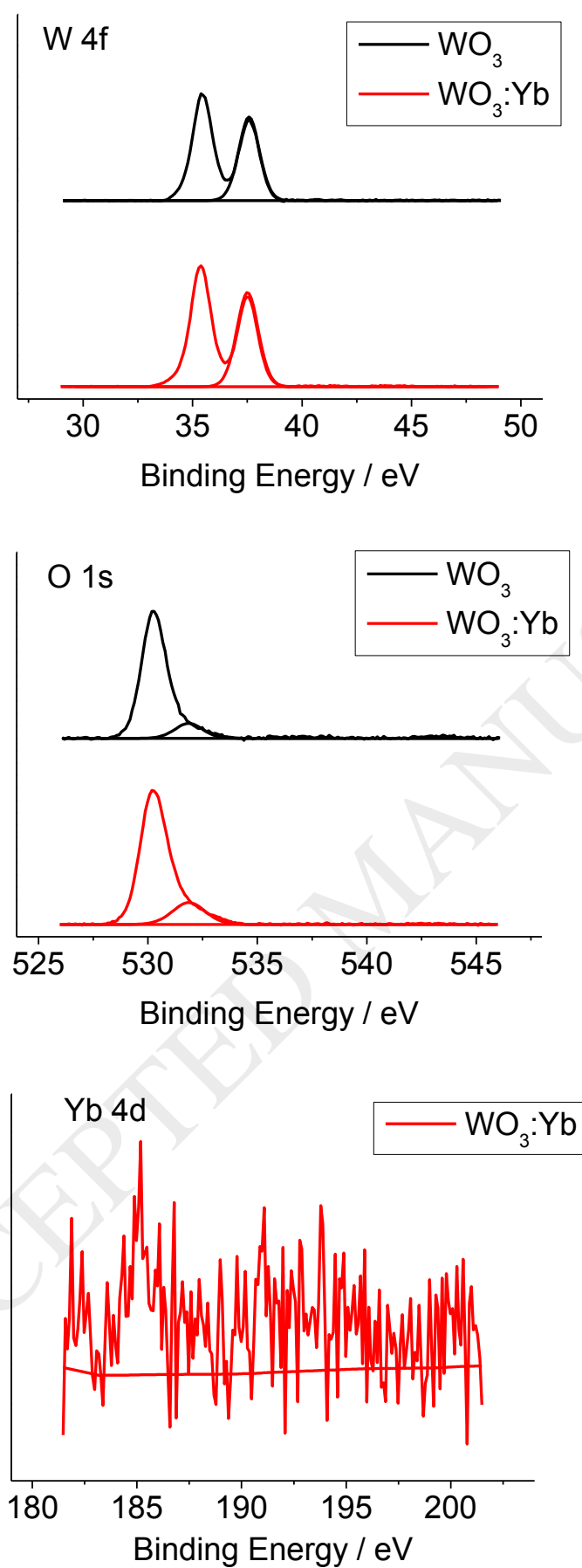
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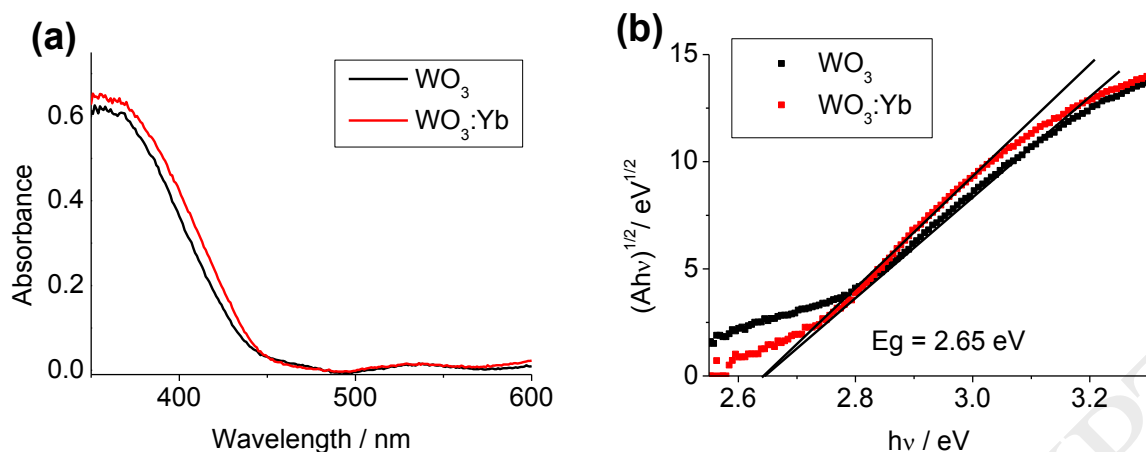
**Figure 1.** XRD patterns for the samples of  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  0.2 at. %.



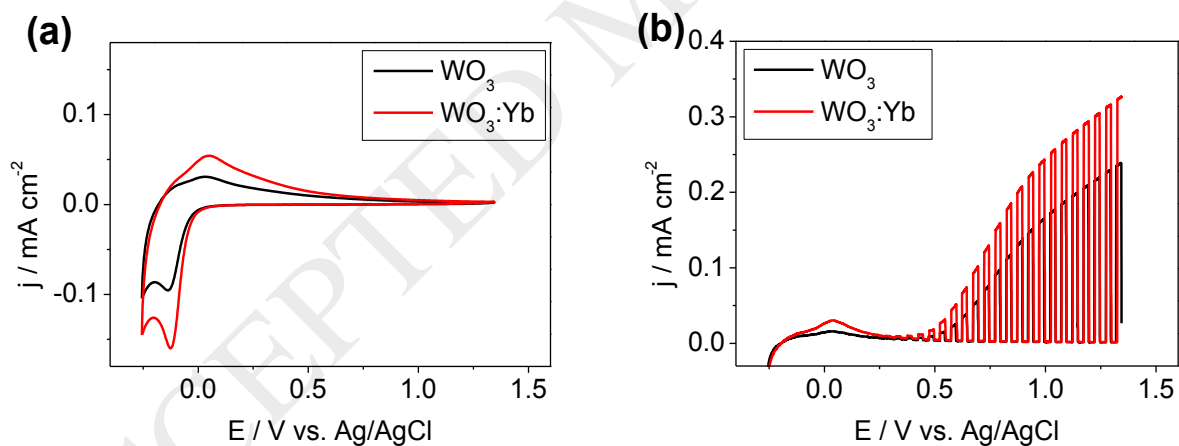
**Figure 2.** Scanning electron micrographs for the samples of (a)  $\text{WO}_3$  and (b)  $\text{WO}_3\text{:Yb 0.2 at\%}$ .



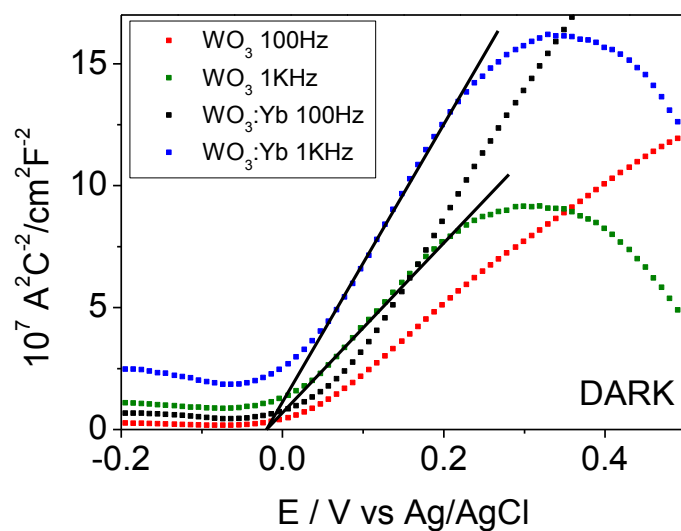
**Fig. 3.** XPS spectra for the samples of  $\text{WO}_3$  and  $\text{WO}_3:\text{Yb}$  0.2 at%.



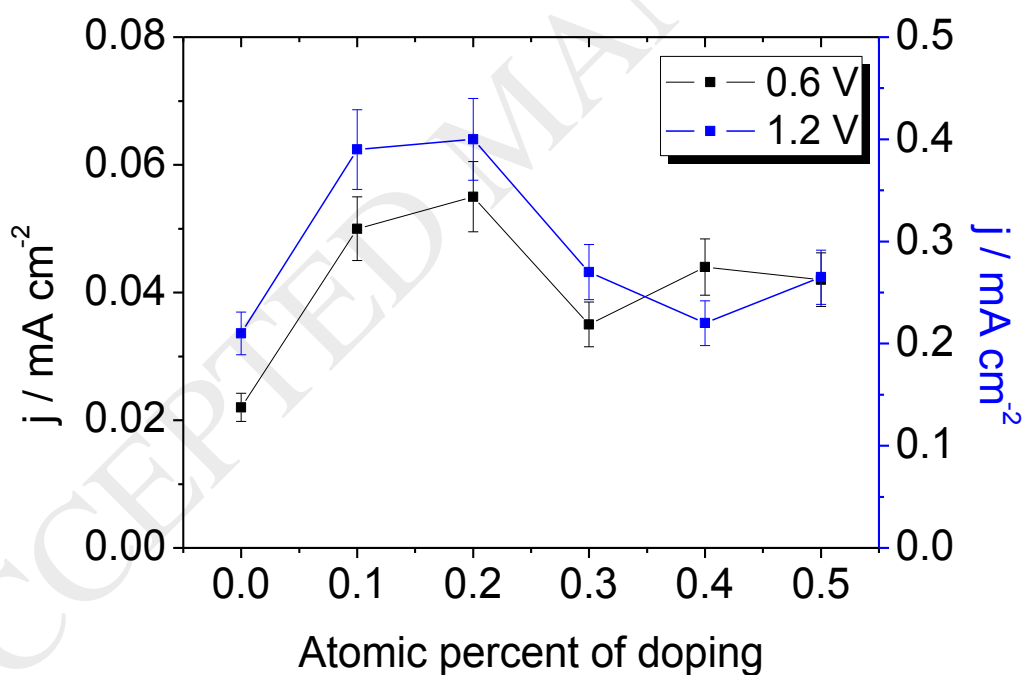
**Fig. 4.** (a) Optical absorbance of the samples of  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  0.2 at.%, (b) Determination of the bandgap from an  $(A\hbar\nu)^{1/2}$  versus  $\hbar\nu$  plot for  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  0.2 at% samples.



**Fig. 5.** (a) Cyclic voltammograms for  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  0.2 at% samples in the dark in 0.1 M  $\text{HClO}_4$ . Scan rate:  $20 \text{ mV s}^{-1}$ . (b) Linear scan voltammograms for the same samples under chopped illumination in 0.1 M  $\text{HClO}_4$ . Scan rate:  $5 \text{ mV s}^{-1}$ .

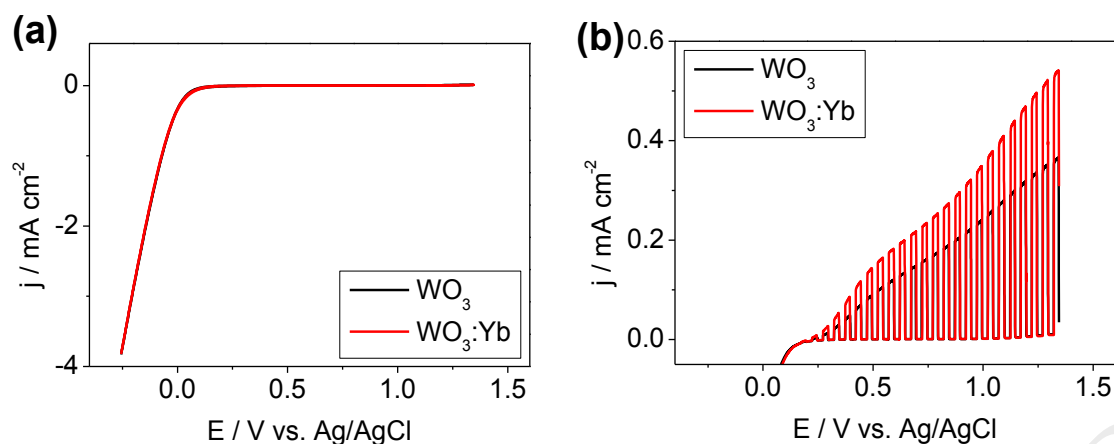


**Fig. 6.** Mott-Schottky plots (at two different frequencies) for  $\text{WO}_3$  and  $\text{WO}_3\text{:Yb}$  0.2 at% samples in 0.1 M  $\text{HClO}_4$  in the dark.

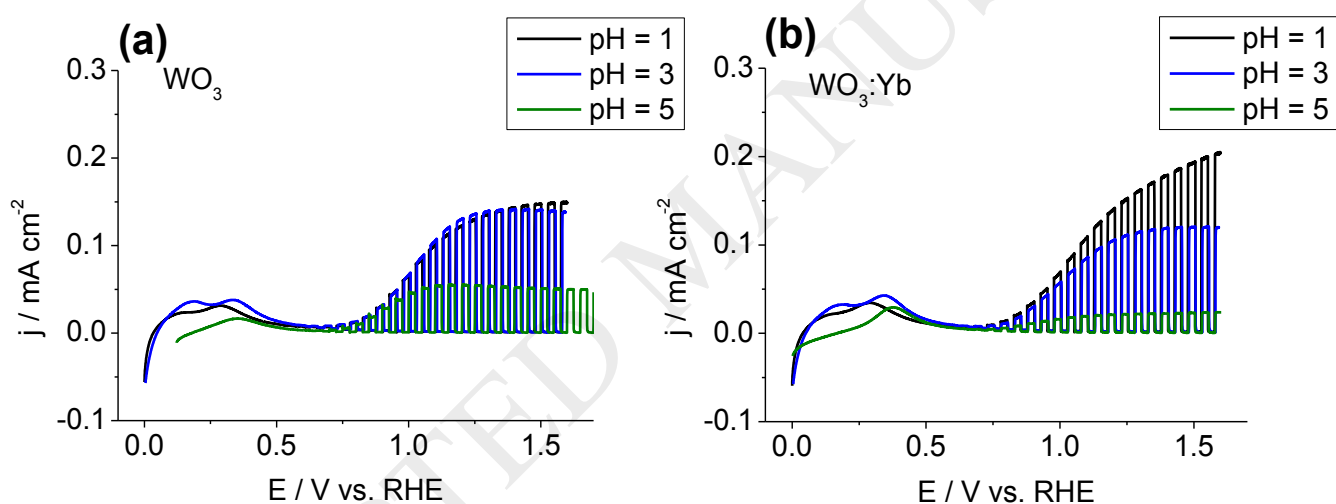


**Fig. 7.** Dependence of the photocurrent density on atomic percent of Yb in the  $\text{WO}_3$  electrodes.





**Fig. 8.** (a) Cyclic voltammograms for  $\text{WO}_3$  and  $\text{WO}_3:\text{Yb}$  0.2 at% samples in the dark in 0.1 M  $\text{HClO}_4$  and  $\text{H}_2\text{O}_2$ . Scan rate:  $20 \text{ mV s}^{-1}$ . (b) Linear scan voltammograms for the samples of  $\text{WO}_3$  and  $\text{WO}_3:\text{Yb}$  0.2 at. % in 0.1 M  $\text{HClO}_4$  and  $\text{H}_2\text{O}_2$ . Scan rate:  $5 \text{ mV s}^{-1}$ .



**Fig. 9.** Linear scan voltammograms under chopped illumination for (a)  $\text{WO}_3$  and (b)  $\text{WO}_3:\text{Yb}$  electrodes in contact with solutions of different pH (1, 3 and 5). Scan rate:  $5 \text{ mV s}^{-1}$ .